nicopure

August 17, 2017

To Whom It May Concern

Statement of Compliance with FDA Establishment Registration and Product Listing Requirements

Dear business partner,

We hereby confirm that Nicopure Labs has complied with the upcoming FDA Establishment Registration and Product Listing Requirements with respect to its manufacturing facilities and its products as follows:

You may find a searchable database of all registered establishments and products at https://www.accessdata.fda.gov/scripts/ctpocerl/index.cfm?action=main.home

Please note that the website path may change from time to time, and that the information may not be always up to date as FDA may be delayed in uploading the information due to the large number of items submitted by various manufacturers.

Under Establishment Registration, Nicopure Labs Manufacturing Gainesville facility was allocated the FEI number 3011318489.

Please also note that we intend to comply with all future requirements with respect to our products as they become effective.

With kind regards,

Patricia Kovacevic

General Counsel, Chief Compliance Officer